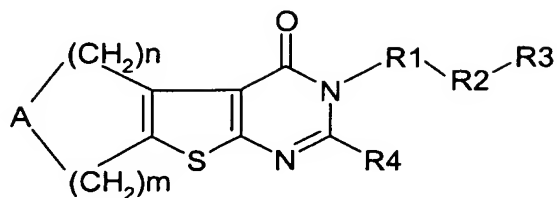


We claim:

1. A compound of the formula (I)



in which

A is O, S, SO, NR⁵ or CH₂;

R⁵ is H, C₁₋₅-alkyl, aryl, aralkyl, acyl or alkoxycarbonyl;

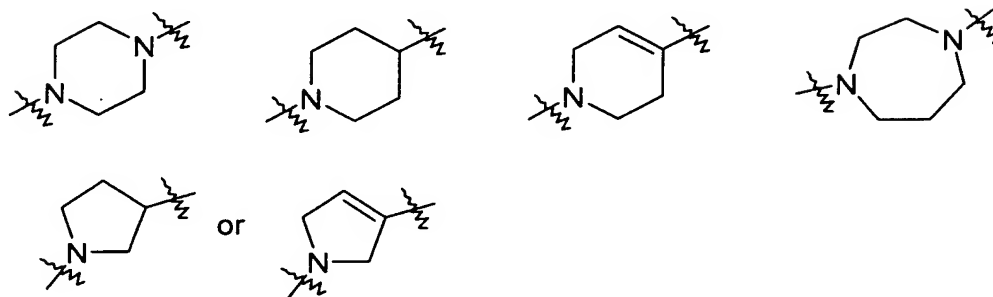
R₄ is H or methyl;

n is 1 or 2;

m is 1 or 2;

R₁ is C₁₋₈-alkylene;

R₂ is a group of the formula



R₃ is 5-membered heteroaryl which may be fused to an aryl or heteroaryl radical, where the heteroaryl and, optionally, the fused aryl or heteroaryl radical may have 1, 2 or 3 substituents selected independently of one another from C₁₋₅-alkyl, C₁₋₅-alkoxy, C₁₋₅-alkylthio, halogen, CN, halo-C₁₋₅-alkyl, halo-C₁₋₅-alkoxy, hydroxy, -NH₂, -N(R₆)₂, -NH(R₆), aryl, aryloxy,

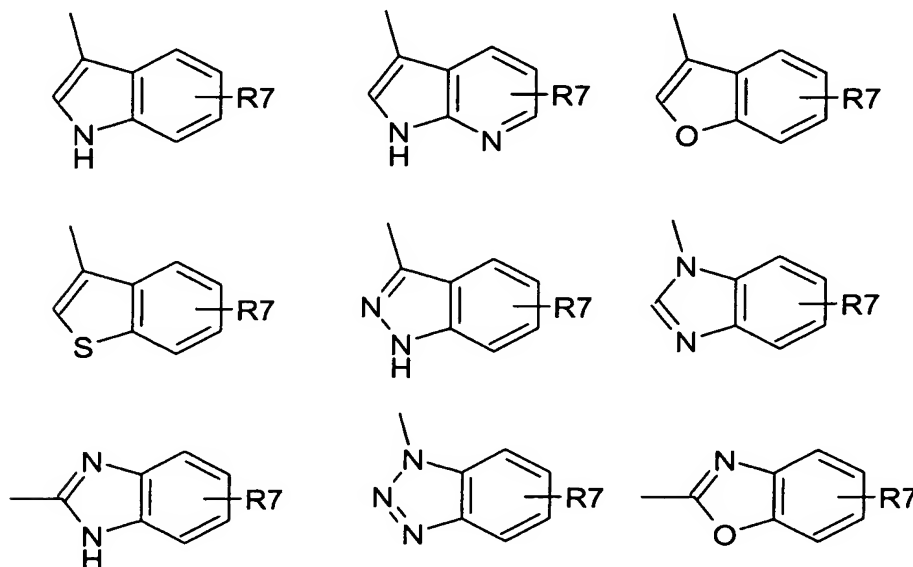
aralkyl, aralkyloxy and heteroaryl, where the substituents aryl, aryloxy, aralkyl, aralkyloxy and heteroaryl may have 1, 2 or 3 substituents selected independently of one another from C₁₋₅-alkyl, C₁₋₅-alkoxy, C₁₋₅-alkylthio, halogen, CN, halo-C₁₋₅-alkyl, halo-C₁₋₅-alkoxy, hydroxy, -NH₂, -N(R₆)₂ and -NH(R₆); and the radicals

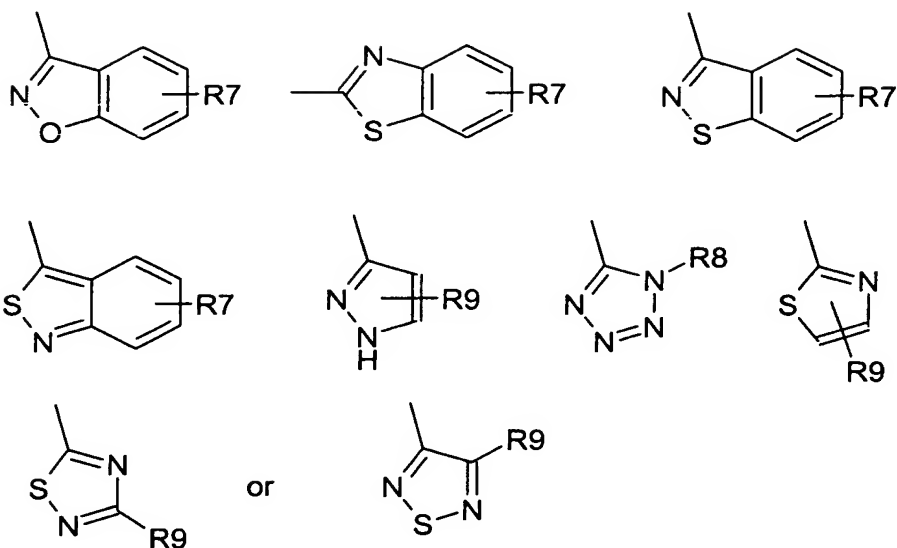
R₆ are independently of one another C₁₋₅-alkyl,

and physiologically tolerated salts thereof.

2. The compound according to claim 1, wherein R₃ is 1H-indol-3-yl, 1H-pyrrolo[2,3-b]pyridin-3-yl, 1-benzofuran-3-yl, 1-benzothien-3-yl, 1H-indazol-3-yl, 1H-benzimidazol-1-yl, 1H-benzimidazol-2-yl, 1H-benzotriazol-1-yl, 1,3-benzoxazol-2-yl, 1,2-benzisoxazol-3-yl, 1,3-benzothiazol-2-yl, 1,2-benzisothiazol-3-yl, pyrazol-3-yl, 1H-tetrazol-5-yl, 1,3-thiazol-2-yl or 1,2,4-thiadiazol-5-yl, which may have 1, 2 or 3 substituents selected independently of one another from C₁₋₅-alkyl, C₁₋₅-alkoxy, halogen, CN, SCH₃, trifluoromethyl, hydroxy, -N(C₁₋₅-alkyl)₂, -NH(C₁₋₅-alkyl), -NH₂, aryl, aryloxy, aralkyl, aralkyloxy and heteroaryl, where the substituents aryl, aryloxy, aralkyl, aralkyloxy and heteroaryl may have 1, 2 or 3 substituents selected independently of one another from C₁₋₅-alkyl, C₁₋₅-alkoxy, halogen, CN, SCH₃, trifluoromethyl, hydroxy, -N(C₁₋₅-alkyl)₂, -NH(C₁₋₅-alkyl) or -NH₂.

3. The compound according to claim 2, wherein R₃ is a radical of the formula

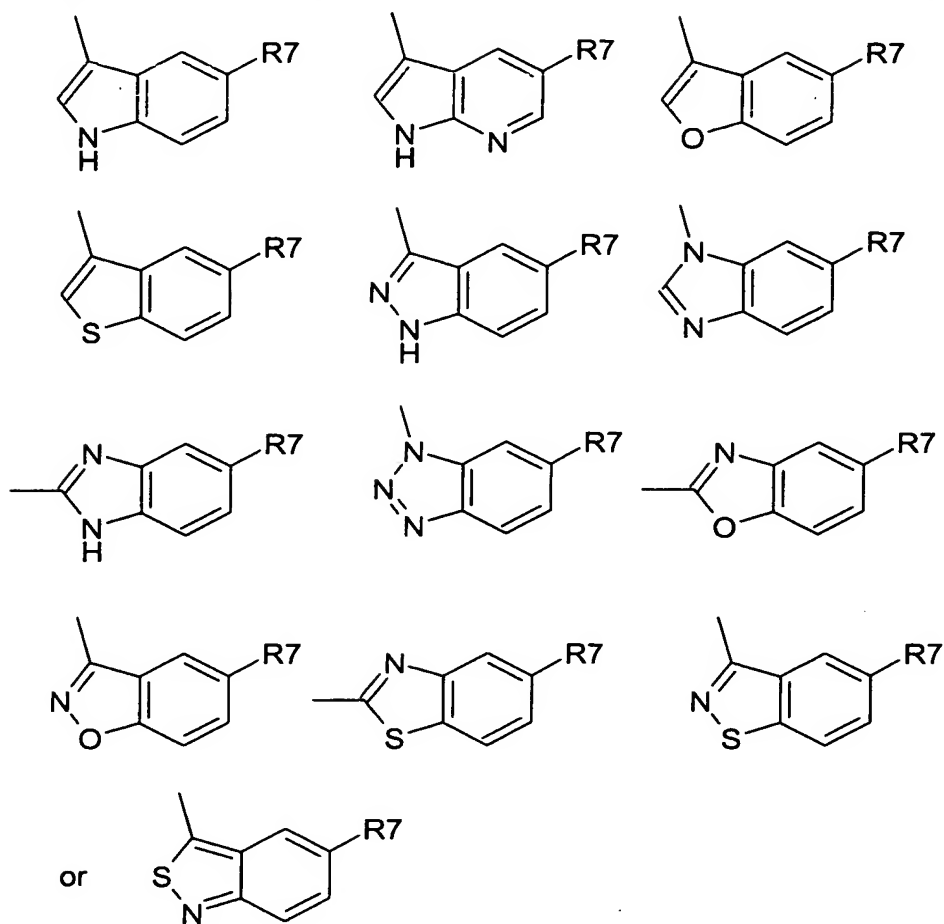




in which

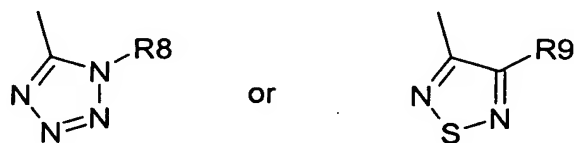
- 5 R7 is H, C₁₋₅-alkyl, C₁₋₅-alkoxy, C₁₋₅-alkylthio, halogen, CN, halo-C₁₋₅-alkyl, halo-C₁₋₅-alkoxy, hydroxy, -NH₂, -N(R6)₂ or -NH(R6); and
- 10 R8 is H, C₁₋₅-alkyl, aryl, aralkyl and heteroaryl;
- 15 R9 is H, C₁₋₅-alkyl, C₁₋₅-alkoxy, C₁₋₅-alkylthio, halogen, CN, halo-C₁₋₅-alkyl, halo-C₁₋₅-alkoxy, hydroxy, -NH₂, -N(R6)₂, -NH(R6), aryl, aryloxy, aralkyl, aralkyloxy or heteroaryl, where aryl, aryloxy, aralkyl, aralkyloxy or heteroaryl may have 1, 2 or 3 substituents selected independently of one another from C₁₋₅-alkyl, C₁₋₅-alkoxy, C₁₋₅-alkylthio, halogen, CN, halo-C₁₋₅-alkyl, halo-C₁₋₅-alkoxy, hydroxy, -NH₂, -N(R6)₂ and -NH(R6); and the radicals
- R6 have the meaning indicated in claim 1.

4. The compound according to claim 3, wherein R3 is a radical of the formula



- 5 in which R7 is as defined in claim 3.

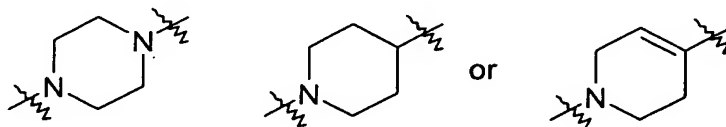
5. The compound according to claim 3, wherein R3 is a radical of the formula



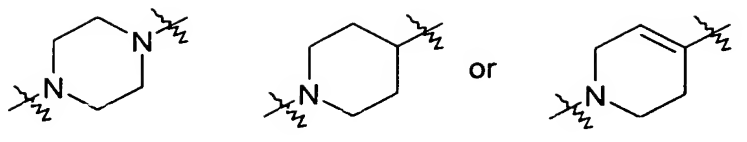
- 10 where R8 and R9 are as defined in claim 3.

6. The compound according to claim 4, wherein R7 is H, C₁₋₅-alkyl, preferably methyl, halogen, preferably chlorine, or halo-C₁₋₅-alkyl, preferably trifluoromethyl.

7. The compound according to claim 5, wherein R8 is C₁₋₅-alkyl, preferably methyl, ethyl or isopropyl or aryl, preferably phenyl.
8. The compound according to claim 5, wherein R9 is C₁₋₅-alkoxy, preferably methoxy, ethoxy or isopropoxy, aryl, preferably phenyl which may be substituted, e.g. by chlorine, or heteroaryl, e.g. 2-thienyl.
9. The compound according to any of claims 1 to 8, wherein A is O, S or NR₅, where R₅ is as defined in claim 1 and is preferably H or methyl.
10. The compound according to any of claims 1 to 8, wherein R4 is hydrogen.
11. The compound according to any of claims 1 to 8, wherein n is 2 and m is 1 or n is 1 and m is 2.
12. The compound according to any of claims 1 to 8, wherein R1 is eth-1,2-ylene, prop-1,3-ylene, prop-1,2-ylene, 2-methyl-prop-1,3-ylene, but-1,2-ylene or but-1,3-ylene.
13. The compound according to any of claims 1 to 8, wherein R2 is a group of the formula



14. The compound according to any of claims 1 to 8, wherein
- R4 is hydrogen;
- n, m are 2, 1 or 1, 2;
- R1 is eth-1,2-ylene, prop-1,3-ylene, prop-1,2-ylene, 2-methylprop-1,3-ylene, but-1,2-ylene or but-1,3-ylene;
- R2 is a group of the formula



and

5 R3 is as defined in any of claims 1 to 13;

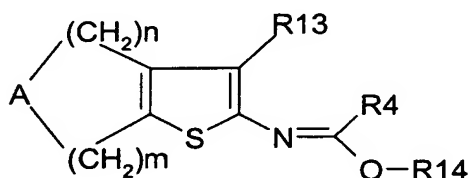
15. The compound according to claim 14, namely
3-substituted 5,6,7,8-tetrahydropyrido[4',3':4,5]thieno[2,3-d]pyrimidin-4(3H)-one
derivatives;

10 3-substituted 3,5,6,8-tetrahydro-4H-pyrano[4',3':4,5]thieno[2,3-d]pyrimidin-4-one
derivatives, or

3-substituted 3,5,6,8-tetrahydro-4H-thiopyrano[4',3';4,5]thieno[2,3-d]pyrimidin-4-
one derivatives.

15 16. A process for preparing a compound according to any of claims 1 to 15

a) by reacting a compound of the formula (II)

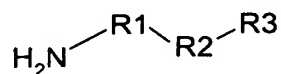


20

in which A, n, m and R4 have one of the meanings indicated in claim 1; R13 is CN
or C₁₋₃-alkyl-O-CO-, and R14 is C₁₋₃-alkyl,

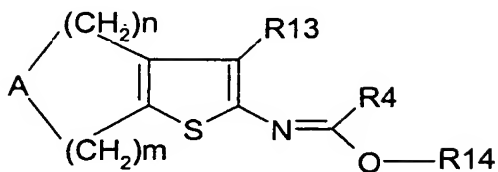
with a primary amine of the formula (III)

25



in which R1, R2 and R3 have one of the meanings indicated in claim 1, and
isolating and, optionally, converting the resulting compound into a physiologically
30 tolerated salt thereof, or

b1) by reacting a compound of the formula (II)



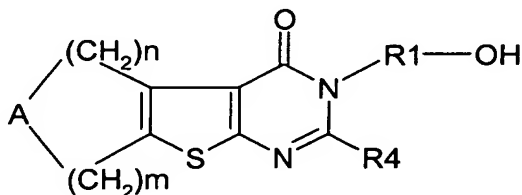
5 in which A, n, m and R_4 have one of the meanings indicated in claim 1; R_{13} is CN or C_{1-3} -alkyl-O-CO-, and R_{14} is C_{1-3} -alkyl,

with a primary amine of the formula (IV)



in which R_1 has one of the meanings indicated in claim 1;

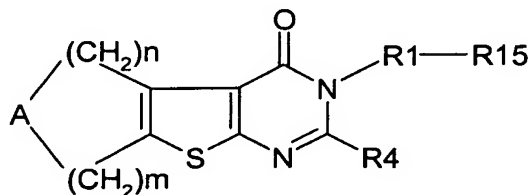
15 b2) reacting the resulting compound of the formula (V)



in which A, n, m, R_4 and R_1 have one of the meanings indicated in claim 1,

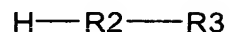
20 with a halogenating agent such as thionyl chloride; and

b3) reacting the resulting compound of the formula (VI)



25 in which A, n, m, R_4 and R_1 have one of the meanings indicated in claim 1, and R_{15} is halogen,

with a secondary amine of the formula (VII)



5

in which R2 and R3 have one of the meanings indicated in claim 1,

and isolating and, optionally, converting the resulting compound into a physiologically tolerated salt thereof.

10

17. The compound according to any of claims 1 to 15 for therapeutic use.

18. A pharmaceutical composition comprising at least one compound according to any of claims 1 to 15 and physiologically acceptable aids.

15

19. The use of a compound according to any of claims 1 to 15 for producing a composition for the treatment of disorders of the central nervous system.

20

20. The use according to claim 19, wherein the disorder of the central nervous system is a neuropsychiatric disorder, in particular a depression.